

INFORMATION DISCLOSURE STATEMENT BY APPLICANT (Not for submission under 37 CFR 1.99)	Application Number		10570229	
	Filing Date		2008-02-28	
	First Named Inventor	Biggadike et al.		
	Art Unit	1624		
	Examiner Name	Berch, Mark L.		
	Attorney Docket Number	PB80476USw		

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	1	0325460	EP					<input type="checkbox"/>
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/Mark Berch/ (02/25/2009)

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4	0741710	EP						<input type="checkbox"/>
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	1	Berge et al., "Pharmaceutical Salts", J. of Pharm. Sci. Vol. 66(1), pp. 1-19, 1977	<input type="checkbox"/>
	2	Daluge et al., "An Efficient, Scalable synthesis of the HIV Reverse Transcriptase Inhibitor Ziagen® (1592U89)", Nucleosides, Nucleotides & Nucleic Acids, Vol. 19, pp. 297-327, 2000	<input type="checkbox"/>
	3	Fleisher et al., "Improved oral drug delivery: solubility limitations overcome by the use of prodrugs", Advanced Drug Delivery Reviews, Vol 19, pp. 115-130, 1996	<input type="checkbox"/>
	4	Legraverend et al., "Synthesis of a New Series of Purine Derivatives and their Anti-Cyclin-Dependent Kinase Activities", J. Heterocyclic Chem., Vol 38, pp. 299-303, 2001	<input type="checkbox"/>

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5	Pederson, et al., "Improved Glucose Tolerance in Zucker Fatty Rats by Oral Administration of the Dipeptidyl Peptidase IV Inhibitor Isoleucine Thiazolidide", Diabetes, Vol. 47, pp. 1253-1258, 1998	<input type="checkbox"/>
6	Vince and Hua, "Synthesis and Anti-HIV Activity of Carbocyclic 2',3'-Didehydro-2',3'-dideoxy 2, 6-Disubstituted Purine Nucleosides" J. Med. Chem., Vol. 33, pp. 17-21, 1990	<input type="checkbox"/>

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